ŞEPA	POTENTIAL HAZARDOU	S WASTE SITE IDEN	TIFICATION	PAN 10167 1576
activity or co be assessed a hazardous v	entification of a potential nfirmation that an actual h under the EPA's Hazardou vaste problem actually exi	nealth or environments is Waste Site Enforcer	al threat exists. A	Il identified sites will
V.S. Alumin	um Lagoons	D. STATE	E. ZIP CODE	FL + BIDGE STS
Marie Ha		PA	17547	Lancaster 071
US Alum		ation .		(17) 424-1781
H. TYPE OF OWNERSHIP (I	(known) 2. STATE 3. COUNTY		S. FRIVATE	6. UNKNOWN
I. SITE DESCRIPTION				
Lagoon	gr basins wh	ich may con	nteun meta	l hydroxide
Audeles.	or basins who The lagoons not he visible	ray be partly	filled in a	r overgrown
and made	I not accusible	as depres	www.	J
	H			
	itizen's complaints, OSHA citat VERNAL - DER 1 AL OR KNOWN PROBLEM			K. DATE IDENTIFIED (mo., day, & yr.) 4/7/80
			21	
Cont	amination of	groundwat	er and S	urface
The Control of the Co	related number	tion and po	aline in	Access no and was
done	tos Minell	anami. 1	Edininary Aluminary	Assessment was u may have
diana	on the	2 . d 1 . c d c . 10	udae Hu	n mag nati-
1 waspag	ed of metal !	garirie se	unge in 1 mg	guary
		F.(*	1	•
a.		PA-208	(
		PA		
				41
M. PREPARER INFORMAT		12	. TELEPHONE RE	G - V FOATE (mo,, day, & yr.)
Lori Da	Uis		7/7 MISSER OF EMER	GEVEDATE (mo., day, & yr.)

EPA Form 2070-8 (5-80)

SEP 1 8 1966

Waste Management



COMMONWEALTH OF PENNSYLVANIA DEPARTMENT OF ENVIRONMENTAL RESOURCES

Post Office Box 2063 Harrisburg, Pennsylvania 17120

ORIGINAL (Red)

Bureau of Waste Management

Preliminary Assessment

FOR

U. S. ALUMINUM LAGOONS PA #2087

> City of Marietta Lancaster County Pennsylvania

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Topography

1/4 Mile Radius

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Section 6 Map of Area Showing Sources of Drinking Water

Section 7 Sources of Information

People Interviewed

Other Sources of Information

Section 9 US Reduction Response Letter To DER Letter in Section 8

 $\frac{\text{Section 10}}{\text{US Reduction Co.}} \quad \text{Analysis of Secondary Aluminum Smelting Waste Products For} \\$

Section 11 Waste Management Assessment for US Reduction Company, Marietta, Pa.

Section 12
Aluminum Recycling Assoc. Work & Report That got the Federal EPA
To Delist The Secondary Aluminum Industry "High Salt Slag" From
The Hazardous Waste Listing.

ORIGINAL (Red)

ORIGINAL (Red)

Narrative / Overview

The site is located at the southwest corner of the Borough of Marietta ONIGINAL at Hazel & Biddle Sts., Lancaster County, Pennsylvania.

The facility is a secondary smelter of aluminum. Wastes generated from this process include air emissions and slag. Prior to 1975, the air emissions were handled by a wet scrubber and the contaminated water went tothe public sewer. U.S. Reduction Company bought U.S. Aluminum in 1975 and installed a baghouse to handle the air emissions; currently the baghouse dust is shipped off site as a hazardous waste containing Cadmium and Lead.

Prior to 1976 the facility operated a slag washing operation to recover aluminum from the slag. Slag from the smelting process was washed with water to dissolve the soluble compoments. The insoluble components were put back into the furnace, The washwater was put into two earthen (unlined) dewatering impoundments; the waterand the soluble components ofthis washwater percolated into the surrounding soil. Insoluble residue would build up inthese impoundments, be periodically dredged, and then be disposed of at an unknown location.

In 1976, the company switched to using a dry recovery operation at another U.S. Reduction plant; the slag was then shipped off site to another plant. U.S. Reduction then closed the two surface impoundments. It is unknown whether or not the impoundments were dredged prior to closure. The impoundments were backfilled and are noe covered over with concrete vy the loading dock. Comapany records (see sections 9,10,11,12) indicate that the slag and slag washwater were not hazardous waste as defined by RCRA. Some of the slag is currently being stored in piles at the southwest corner of the plant; off site recovery operation had shut down for a while.

The Borough of Marietta has received complaint from nearby residents that that emissions from this plant are corroding the paint off their houses. This may indicate problems with the air polution control equipment used by the facility and possible air, soil, and surface water contamination with Cadmiun and lead.

ORIGINAL (Red)

The low assessment was not based on the lagoons since company records indicate that waste in lagoons was not a RCRA defined hazardous waste; the low assessment was based onthe alleged air emissions and possible air, soil, and surface water contamination.

Geology / Hydrology

The site is located in the floodplain of the Susquehanna River at the toe of a moderately steep slope from the north; the area of the facility itself is relatively flat. The area from Market St. slopes down to flat area where the lagoons were. The built up railroad bed on the south side of the facility acts as a dike/berm between the site and the River.

ORIGINAL

The site is underlain by an undetermined depth of sandy alluvium. The bedrock under the site is of the Vintage Formation. This is a dolomite formation with groundwater flow through solution channels.

Drinking Water

Drinking water for the area comes from privite wells and the Marietta Gravity Water Company. The Marietta Gravity Water Company services the borough of Marietta and a few large buildings north of Marietta on the south side of Route 441. The Marietta Gravity Water Company has the following sources of water:

- 3 groundwater wells located on west side of Route 441 in the village of Chickies, Lancaster County.
- Two reservoirs in Helem Township, York County which are fed by 52 springs in that area. The water from these reservoirs ispiped under the Susquehanna River.

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(hed)

Section 2

,



POTENTIAL HAZARDOUS WASTE SITE

I. IDENT	IFICATION
O1 STATE PA	2087

	PRELIMINARY SITE INFORMA			1ENT	2087
II. SITE NAME AND LOCATION					
01 SITE NAME (Legal, common, or descriptive name of site)		02 STREET	. ROUTE NO. O	R SPECIFIC LOCATION IDENTIFIE	
U. S. Aluminum Lagoons		1950105	1 & Bidd		10
Marietta		Pa.	05 ZIP CODE 17547	OG COUNTY Lancaster	07COUNTY 08 CONG COPE DIST 07 16
09 COORDINATES LATITUDE LONG 40 32 54" 076 3:	3 30°.				10.1 110
Site located at south west cornerstreet intersects with Route 44	er of Mari	etta Bo	orough o rough.	ff Market Street	. Market
III. RESPONSIBLE PARTIES					
01 OWNER (// known)		02 STREET	(Business, mailing, i	residentiali	
U.S. Reduction Co.		2025			
03 CITY			05 ZIP CODE	06 TELEPHONE NUMBER	T
Lansing		III.	50438	800)323-8760	
07 OPERATOR (if known and different from owner)		08 STREET	(Business, mailing, r	esidential)	
U.S. Aluminum Corp.		Haze	l & Bidd	le Sts.	
09 CITY			1 ZIP CODE	12 TELEPHONE NUMBER	
Marietta		Pa.	17547	(717) 426-1981	
13 TYPE OF OWNERSHIP (Check one)		1 4.	17547	717.420-1301	
💢 A. PRIVATE 🗆 B. FEDERAL:	(Agency name)		C. STAT	E □D.COUNTY □ E. N	IUNICIPAL
☐ F. OTHER:	(Agency name)		☐ G. UNKN		
(Specity) 14 OWNER/OPERATOR NOTIFICATION ON FILE (Check all that apply)			L G. ONK	NOWIN .	
A. RCRA 3001 DATE RECEIVED: / / MONTH DAY YEAR	B UNCONTROLL	ED WASTE	CITE		- //
IV. CHARACTERIZATION OF POTENTIAL HAZARD	B. GIACOIATHOLL	EDWASIE	SITE (CERCLA 10)	DATE RECEIVED:	DAY YEAR C. NONE
01 ON SITE INSPECTION					
XYES DATE 3 ,19 , 87	all that apply) A 🔲 B. EPA CAL HEALTH OFFIC	CONTRAC	TOR X		RCONTRACTOR
CONTRA	CTOR NAME(S): _			(Specify)	
	03 YEARS OF OPERA	TION	15		
□ A. ACTIVE □ B. INACTIVE □ C. UNKNOWN 04 DESCRIPTION OF SUBSTANCES POSSIBLY PRESENT, KNOWN, OF	BE	GINNING YEAR	ENDING	YEAR UNKNOW	VN
		+:1 01	osupo i	. 1076	
Lagoons were operated from approindicate that waste that went in Cd & Pb has been found in baghou	ito lagoons ise dust.	was r	not hazan	rdous as defined	records by RCRA.
os description of potential HAZARD TO ENVIRONMENT AND/OR Prior to 1975 the furnace emissi went to public sewer; in 1975 th neighbors that air emissions fro	ons were h le baghouse	was i	nstalled	d. Complaints re	the water ceived form
V. PRIORITY ASSESSMENT					
D1 PRIORITY FOR INSPECTION (Check one. If high or medium is checked, comp	elete Part 2 - Waste Inform.	ation and Part 3	- Description of Haza	ordous Conditions and Incidents	
☐ A. HIGH [Inspection required promptly] ☐ B. MEDIUM [Inspection required]	C. LOW (Inspect on time as		D. NONE		sition form)
VI. INFORMATION AVAILABLE FROM					
1 CONTACT 0	2 OF (Agency/Organizati	on)			03 TELEPHONE NUMBER
Thomas Sheehan, Project Officer	Pa. DER, Div	. Emerg	ency & Rem	edial Response	215 3565-1687
04 PERSON RESPONSIBLE FOR ASSESSMENT 0	5 AGENCY	06 ORGANIZ		07 TELEPHONE NUMBER	08 DATE
Thomas Sheehan, Project Officer	Pa. DER	Waste	Manageme	ent (215)565-1687	4 ,27,87 MONTH DAY YEAR



		fundal	
ı.	IDEN	TIFICATION	

Ş E		PO	TENTIAL HAZA	RDOUS WAS	TE SITE		ICATION
			PRELIMINAR PART 2 - WAS	Y ASSESSMENTE INFORMATION	NT	01 STATE 0	2 SITE NUMBER
I. WASTES	STATES, QUANTITIES, A	ND CHARACTE	RISTICS				
01 PHYSICAL:	STATES (Check all that apply)	02 WASTE QUAN		03 WASTE CHARA	CTERISTICS (Check all that	apply)	
X A. SOLID X B. POWD	☐ E. SLURRY	must	e independent)	XA. TOX	IC □ E SOL		IGHLY VOLATILE
C. SLUDG	ER, FINES F, LIQUID E XG, GAS	TONS		☐ B. COR	ROSIVE I F. INFE	CTIOUS	XPLOSIVE
☐ D. OTHER	0.5 0 € 05455000000	CUBIC YARDS		D. PER		TABLE 🗀 L. I	REACTIVE NCOMPATIBLE
	(Specify)	NO. OF DRUMS				□ M.	NOT APPLICABLE
II. WASTE 1	YPE						
CATEGORY	SUBSTANCE N	IAME	01 GROSS AMOUNT	02 UNIT OF MEASU	DE 03 COLUMN		
SLU	SLUDGE			SE SITT OF MEASO	RE 03 COMMENTS		
OLW	OILY WASTE						
SOL	SOLVENTS						
PSD	PESTICIDES			-			
осс	OTHER ORGANIC CH	HEMICALS					
IOC	INORGANIC CHEMIC						
ACD	ACIDS				25 Page 1		
BAS	' BASES						
MES	HEAVY METALS						
. HAZARDO	OUS SUBSTANCES (See AL			<u> </u>	Cd & Pb ir	n baghouse	dust
CATEGORY	02 SUBSTANCE N						
MES	Cadmium	AME	03 CAS NUMBER		SPOSAL METHOD	05 CONCENTRAT	ION 06 MEASURE (
MES	Lead		7440-43-9	Baghouse/	off site		
120	Leau			Baghouse/	off site		
	The second of the second						
FEDET							
	KS (See Appendix for CAS Numbers	NA					
CATEGORY	01 FEEDSTOCK	NAME	02 CAS NUMBER	CATEGORY	01 FEEDSTO	CK NAME	02.040.4
FDS			2000	FDS			02 CAS NUMBER
FDS				FDS			
FDS		9		FDS		101	
FDS							
OURCES	OF INFORMATION (Cité spe	cific references & a	No blas service	FDS			
	Crima I IOII leve spe	concrete ences, e.g., st.	ate riles, sample analysis, rep	ports)	7		
	58 HH#0	1 1					
See	se t tion 7.	1				30	
	NO 17.4500 15.500 N 18 18 18 18 18 18 18 18 18 18 18 18 18 18				00		
		2 22					

ONIGINAL (Red)

S.EDA

POTENTIAL HAZARDOUS WASTE SITE

I. IDENTIFICATION

01 STATE 02 SITE NUMBE

WEIA	PR PART 3 - DESCRIPTION	ELIMINARY ASSESSMENT OF HAZARDOUS CONDITIONS AND	INCIDENTO	O1 STATE O	2087
II. HAZARDOUS CONDITIO	NS AND INCIDENTS	AND CONDITIONS AND	INCIDENTS		
01 A. GROUNDWATER C	ONTAMINATION	02 G OBSERVED (DATE:) DF	POTENTIAL	ALLEGED
Groundwater cont	amination with	O2 OBSERVED (DATE: O4 NARRATIVE DESCRIPTION residual waste material ; Company records indica	1		
waste is not haz emissions; parti	ardous as define	o2 OBSERVED (DATE: 04 NARRATIVE DESCRIPTION on the ground; Company ed by RCRA. Complaints re ay be contaminating surf	records ind	otential licate t arding	▼ALLEGED that this air
THE TECTE BOT OUGH	has received or	02 © OBSERVED (DATE: 04 NARRATIVE DESCRIPTION Omplaints from nearby re: nt off their houses.	sidences th	otential at emis	⋉ ALLEGED
01 D. FIRE/EXPLOSIVE CO	NDITIONS	02 C OBSERVED (DATE:			
None found by th	is investigation	04 NAHHATIVE DESCRIPTION		TENTIAL	□ ALLEGED
01 [E. DIRECT CONTACT 03 POPULATION POTENTIALL None found by thi) 🗆 PO	TENTIAL	□ ALLEGED
D1 S F. CONTAMINATION OF D3 AREA POTENTIALLY AFFEC	SOIL 2	02 G OBSERVED (DATE.			
esidual waste st hat this waste w rom air emission	ored in earthed ea not hazardous s (Cd & Pb).	o4 NARRATIVE DESCRIPTION dewatering impoundments s as defined by RCRA. Pos		ecords contam	□ ALLEGED indicate ination
1 KG. DRINKING WATER CON 3 POPULATION POTENTIALLY		02 OBSERVED (DATE:) X POT	ENTIAL	
ancaster Water A	contamination ha uthority surface	o4 NARRATIVE DESCRIPTION as happened the Columbia e intakes downstream on t	200		□ ALLEGED y of ver may
 H. WORKER EXPOSURE/I WORKERS POTENTIALLY A 	NJURY	02 OBSERVED (DATE:)	The state of the s	
ne found by this		04 NARRATIVE DESCRIPTION		ENTIAL	□ ALLEGED
I. POPULATION EXPOSURE	INJURY	02 🗆 OBSERVED (DATE:	7 1 7 200-		
rietta borough h	TECTED:	02 OBSERVED (DATE: 04 NARRATIVE DESCRIPTION Dlaints from nearby resident, off their houses) □ POTE	NTIAL }	ALLEGED

ORIGINAL

SEDA

POTENTIAL HAZARDOUS WASTE SITE

I. IDENTIFICATION

PRE	LIMINARY ASSESSMENT .	01 STATE 02 SITE NUMBER
PART 3 - DESCRIPTION	OF HAZARDOUS CONDITIONS AND INCIDENTS	PA 2087
II. HAZARDOUS CONDITIONS AND INCIDENTS (Continu	ed)	
01 □ J. DAMAGE TO FLORA 04 NARRATIVE DESCRIPTION	02 GBSERVED (DATE:)	
STANTIANTE DESCRIPTION		□ POTENTIAL □ ALLEGED
None found by this investiga		(**)
Hone round by this investiga	cion.	
01 7 8 20005 70 70		
01 ☐ K. DAMAGE TO FAUNA 04 NARRATIVE DESCRIPTION (Include name(s) of species)	02 - OBSERVED (DATE:)	□ POTENTIAL □ ALLEGED
593		☐ POTENTIAL ☐ ALLEGED
None found by this investiga	tion.	
- 397		
01 □ L. CONTAMINATION OF FOOD CHAIN	00 17 00000	
04 NARRATIVE DESCRIPTION	02 OBSERVED (DATE:)	POTENTIAL ALLEGED
None found by this investice	1.5	
None found by this investiga	tion.	
01 M. UNSTABLE CONTAINMENT OF WASTES	03 D OBSERVED IN LTD	
(Spills/runoff/standing liquids/feaking drums) 03 POPULATION POTENTIALLY AFFECTED:	02 OBSERVED (DATE:)	POTENTIAL ALLEGED
Residual waste stored in miles	— 04 NARRATIVE DESCRIPTION	
Residual waste stored in piles; chazardous as defined by RCRA.	company records indicate that thi	is waste is not
TTTTTTT BY NOWN.		
01 M. DAMAGE TO OFFSITE PROPERTY 04 NARRATIVE DESCRIPTION	02 OBSERVED (DATE:)	POTENTIAL MALLEGED
Marietta Borough has received con	anlaint. Co	POTENTIAL MALLEGED
Marietta Borough has received con from the plant are corroding the	praints from nearbt residences t	chat air emissions
and praise and corroding the	paint off their houses.	
01 C O CONTAMINATION OF SEWERS AND		
01 \square 0. CONTAMINATION OF SEWERS, STORM DRAINS, W 04 NARRATIVE DESCRIPTION	WTPs 02 G OBSERVED (DATE:)	POTENTIAL ALLEGED
None found by this investigati		
mene round by this investigati	011.	
01 D. ILLEGAL/UNAUTHORIZED DUMPING	02 - OBSERVED (DATE	
U4 NARRATIVE DESCRIPTION		POTENTIAL ALLEGED
None found by this investigati	on.	
9		
		#
05 DESCRIPTION OF ANY OTHER KNOWN, POTENTIAL, OR A	LLEGED HAZARDS	
Alleged air emissions may contain	Cd and Ph. may contaminate sunfi	200 112102 :1
or be inhaled directly.	and is, may containfinate suits	ace water, soil,
TOTAL POPULATION POTENTIAL		
. TOTAL POPULATION POTENTIALLY AFFECTED: COMMENTS		
Maste handled in the unlined surfaceRA) according to company records	ace impoundments was no hazardous	las defined bu
RCRA) according to company records		, tas der med by
es P		
SOURCES OF INFORMATION		
SOURCES OF INFORMATION (Cite specific references, e.g., state to	iles, sample analysis, reports)	
See sections 7,8,9,10,11,12,13.		
1,0,0,0,10,11,12,13.		

APPENDIX

ORIGINAL (Red)

I. FEEDSTOCKS

CAS Number	Chemical Name		*		
or to trainbur	Chemical Name	CAS Number	Chemical Name	CAS Number	Chemical Name
1. 7664-41-7 2. 7440-36-0 3. 1309-64-4 4. 7440-38-2 5. 1327-53-3 6. 21109-95-5 7. 7726-95-6 8. 106-99-0 9. 7440-43-9	Ammonia Antimony Antimony Trioxide Arsenic Arsenic Trioxide Barium Sulfide Bromine Butadiene Cadmium	CAS Number 14. 1317-38-0 15. 7758-98-7 16. 1317-39-1 17. 74-85-1 18. 7647-01-0 19. 7664-39-3 20. 1335-25-7 21. 7439-97-6	Chemical Name Cupric Oxide Cupric Sulfate Cuprous Oxide Ethylene Hydrochloric Acid Hydrogen Fluoride Lead Oxide Mercury	27. 7778-50-9 28. 1310-58-3 29. 115-07-1 30. 10588-01-9 31. 1310-73-2 32. 7646-78-8 33. 7772-99-8 34. 7664-93-9	Chemical Name Potassium Dichromate Potassium Hydroxide Propylene Sodium Dichromate Sodium Hydroxide Stannic Chloride Stannous Chloride Sulfuric Acid
10. 7782-50-5 11. 12737-27-8 12. 7440-47-3 13. 7440-48-4	Chlorine Chromite Chromium Cobalt	22. 74-82-8 23. 91-20-3 24. 7440-02-0 25. 7697-37-2 26. 7723-14-0	Methane Napthalene Nickel Nitric Acid Phosphorus	35. 108-88-3 36. 1330-20-7 37. 7646-85-7 38. 7733-02-0	Toluene Xylene Zinc Chloride Zinc Sulfate

II. HAZARDOUS SUBSTANCES

CAS Number	Chemical Name				
	Chemical Name	CAS Number	Chemical Name	CAS Number	Chemical Name
1. 75-07-0	Acetaldehyde	47. 1303-33-9	Arsenic Trisulfide	92. 142-71-2	
2. 64-19-7	Acetic Acid	48. 542-62-1	Barium Cyanide	93. 12002-03-8	Cupric Acetate
3. 108-24-7	Acetic Anhydride	49. 71-43-2	Benzene	94. 7447-39-4	
4. 75-86-5	Acetone Cyanohydrin	50. 65-85-0	Benzoic Acid	95. 3251-23-8	Cupric Chloride
5. 506-96-7	Acetyl Bromide	51. 100-47-0	Benzonitrile		Cupric Nitrate
6. 75-36-5	Acetyl Chloride	52. 98-88-4	Benzoyl Chloride	96. 5893-66-3	Cupric Oxalate
7. 107-02-8	Acrolein	53. 100-44-7	Benzyl Chloride	97. 7758-98-7	Cupric Sulfate
8. 107-13-1	Acrylonitrile	54. 7440-41-7	Beryllium	98. 10380-29-7	re
9. 124-04-9	Adipic Acid	55. 7787-47-5	Beryllium Chloride	99. 815-82-7	Cupric Tartrate
10. 309-00-2	Aldrin	56. 7787-49-7	Beryllium Fluoride	100. 506-77-4	Cyanogen Chloride
11. 10043-01-3	Aluminum Sulfate	57. 13597-99-4	Beryllium Nitrate	101.110-82-7	Cyclohexane
12. 107-18-6	Allyl Alcohol	58. 123-86-4	Butyl Acetate	102.94-75-7	2,4-D Acid
13. 107-05-1	Allyl Chloride	59. 84-74-2	n-Butyl Phthalate	103. 94-11-1	2,4-D Esters
14. 7664-41-7	Ammonia	60. 109-73-9	Butylamine	104. 50-29-3	DDT
15. 631-61-8	Ammonium Acetate	61, 107-92-6	Butyric Acid	105. 333-41-5	Diazinon
16. 1863-63-4	Ammonium Benzoate	62. 543-90-8	Cadimium Acetate	106. 1918-00-9	Dicamba
17. 1066-33-7	Ammonium Bicarbonate	63. 7789-42-6	Cadmium Bromide	107. 1194-65-6	Dichlobenil
18. 7789-09-5	Ammonium Bichromate	64. 10108-64-2	Cadmium Chloride	108. 117-80-6	Dichlone
19. 1341-49-7	Ammonium Bifluoride	65. 7778-44-1	Calcium Arsenate	109. 25321-22-6	Dichlorobenzene (all isomers)
20. 10192-30-0	Ammonium Bisulfite	66. 52740-16-6	Calcium Arsenite	110. 266-38-19-7	Dichloropropane (all isomers)
21. 1111-78-0	Ammonium Carbamate	67. 75-20-7	Calcium Carbide	111. 26952-23-8	Dichloropropene (all isomers)
22. 12125-02-9	Ammonium Chloride	68. 13765-19-0	Calcium Chromate	112.8003-19-8	Dichloropropene-
23. 7788-98-9	Ammonium Chromate	69. 592-01-8	Calcium Cyanide	110 75 00 0	Dichloropropane Mixture
24. 3012-65-5	Ammonium Citrate, Dibasic	70. 26264-06-2	Calcium Dodecylbenzene	113. 75-99-0	2-2-Dichloropropionic Acid
25. 13826-83-0	Ammonium Fluoborate		Sulfonate	114.62-73-7	Dichlorvos
26. 12125-01-8	Ammonium Fluoride	71, 7778-54-3	Calcium Hypochlorite	115.60-57-1	Dieldrin
27. 1336-21-6	Ammonium Hydroxide	72. 133-06-2	Captan	116. 109-89-7 117. 124-40-3	Diethylamine
28.6009-70-7	Ammonium Oxalate	73. 63-25-2	Carbaryl		Dimethylamine
29. 16919-19-0	Ammonium Silicofluoride	74. 1563-66-2	Carbofuran	118. 25154-54-5 119. 51-28-5	Dinitrobenzene (all isomers)
30. 7773-06-0	Ammonium Sulfamate	75. 75-15-0	Carbon Disulfide	120, 25321-14-6	Dinitrophenol
31. 12135-76-1	Ammonium Sulfide	76. 56-23-5	Carbon Tetrachloride	121. 85-00-7	Dinitrotoluene (all isomers)
32. 10196-04-0	Ammonium Sulfite	77. 57-74-9	Chlordane	122. 298-04-4	Diquat
33. 14307-43-8	Ammonium Tartrate	78. 7782-50-5	Chlorine	123. 330-54-1	Disulfoton
34. 1762-95-4	Ammonium Thiocyanate	79. 108-90-7	Chlorobenzene	124. 27176-87-0	Diuron
35. 7783-18-8	Ammonium Thiosulfate	80. 67-66-3	Chloroform	125. 115-29-7	Dodecylbenzenesulfonic Acid
36. 628-63-7	Amyl Acetate	81.7790-94-5	Chlorosulfonic Acid	126. 72-20-8	Endosulfan (all isomers)
37. 62-53-3	Aniline	82. 2921-88-2	Chlorpyrifos	127. 106-89-8	Endrin and Metabolites
38. 7647-18-9	Antimony Pentachloride	83. 1066-30-4	Chromic Acetate	128. 563-12-2	Epichlorohydrin
39. 7789-61-9	Antimony Tribromide	84. 7738-94-5	Chromic Acid	129. 100-41-4	Ethion
40. 10025-91-9	Antimony Trichloride	85. 10101-53-8	Chromic Sulfate	130, 107-15-3	Ethyl Benzene
41. 7783-56-4	Antimony Trifluoride	86. 10049-05-5	Chromous Chloride	131. 106-93-4	Ethylenediamine .
42. 1309-64-4	Antimony Trioxide	87. 544-18-3	Cobaltous Formate	132, 107-06-2	Ethylene Dibromide
43. 1303-32-8	Arsenic Disulfide	88. 14017-41-5	Cobaltous Sulfamate	133, 60-00-4	Ethylene Dichloride
44. 1303-28-2	Arsenic Pentoxide	89. 56-72-4	Coumaphos	134, 1185-57-5	EDTA
45. 7784-34-1	Arsenic Trichloride	90. 1319-77-3	Cresol	135, 2944-67-4	Ferric Ammonium Citrate
46. 1327-53-3	Arsenic Trioxide	91.4170-30-3	Crotonaldehyde	136. 7705-08-0	Ferric Ammonium Oxalate
					Ferric Chloride

II. HAZARDOUS SUBSTANCES

- 30-301-354 - 31-305-35and the

1 3 . . 104

	II. HAZARDO	US SUBSTANCES				ORIGINAL
	CAS Number	Chemical Name	CAS Number	Chemical Name	CAS Number	Chemical Name
	137. 7783-50-8	Ferric Fluoride	192, 74-89-5	Monomethylamine	249. 7632-00-0	Sodium Nitrate
	138. 10421-48-4	Ferric Nitrate	193. 300-76-5	Naled '	250. 7558-79-4	Sodium Phosphate, Dibasic
	139. 10028-22-5	Ferric Sulfate	194, 91-20-3	Naphthalene	251. 7601-54-9	Sodium Phosphate, Tribasic
	140. 10045-89-3	Ferrous Ammonium Sulfate	195. 1338-24-5	Naphthenic Acid	252, 10102-18-8	
٠.	141.7758-94-3	Ferrous Chloride	196. 7440-02-0	Nickel	253. 7789-06-2	Strontium Chromate
	142. 7720-78-7	Ferrous Sulfate	197, 15699-18-0	Nickel Ammonium Sulfate	254. 57-24-9	Strychnine and Salts
	143. 206-44-0	Fluoranthene	198. 37211-05-5	Nickel Chloride	255. 100-420-5	Styrene
	144. 50-00-0	Formaldehyde	199. 12054-48-7	Nickel Hydroxide	256. 12771-08-3	
	145.64-18-6	Formic Acid	200. 14216-75-2	Nickel Nitrate	257. 7664-93-9	Sulfuric Acid
	146. 110-17-8	Fumaric Acid	201. 7786-81-4	Nickel Sulfate	258. 93-76-5	2,4,5-T Acid
	147. 98-01-1 148. 86-50-0	Furfural Guthion	202. 7697-37-2	Nitric Acid	259. 2008-46-0	2,4,5-T Amines
	149. 76-44-8	Heptachlor	203. 98-95-3	Nitrobenzene	260. 93-79-8	2,4,5-T Esters
	150. 118-74-1	Hexachlorobenzene	204. 10102-44-0	Nitrogen Dioxide	261. 13560-99-1	
	151. 87-68-3	Hexachlorobutadiene	205. 25154-55-6	Nitrophenol (all isomers)	262.93-72-1	2,4,5-TP Acid
	152. 67-72-1	Hexachloroethane	206. 1321-12-6	Nitrotoluene	263. 32534-95-5	
	153. 70-30-4	Hexachlorophene	207. 30525-89-4 208. 56-38-2	Paraformaldehyde	264. 72-54-8	TDE
	154. 77-47-4	Hexachlorocyclopentadiene	209. 608-93-5	Parathion	265. 95-94-3	Tetrachlorobenzene
	155. 7647-01-0	Hydrochloric Acid	210. 87-86-5	Pentachlorobenzene	266. 127-18-4	Tetrachloroethane
		(Hydrogen Chloride)	211. 85-01-8	Pentachlorophenol Phenanthrene	267. 78-00-2	Tetraethyl Lead
	156. 7664-39-3	Hydrofluoric Acid	212, 108-95-2	Phenoi	268. 107-49-3 269. 7446-18-6	Tetraethyl Pyrophosphate Thallium (I) Sulfate
		(Hydrogen Fluoride)	213. 75-44-5	Phosgene	270. 108-88-3	Toluene
	157. 74-90-8 "	Hydrogen Cyanide	214. 7664-38-2	Phosphoric Acid	271.8001-35-2	Toxaphene
	158. 7783-06-4	Hydrogen Sulfide	215, 7723-14-0	Phosphorus	272. 12002-48-1	Trichlorobenzene (all isomers)
	159. 78-79-5	Isoprene	216. 10025-87-3	Phosphorus Oxychloride	273. 52-68-6	Trichlorfon
	160. 42504-46-1	Isopropanolamine	217. 1314-80-3	Phosphorus Pentasulfide	274. 25323-89-1	Trichloroethane (all isomers)
		Dodecylbenzenesulfonate	218. 7719-12-2	Phosphorus Trichloride	275. 79-01-6	Trichloroethylene
	161. 115-32-2	Kelthane	219. 7784-41-0	Potassium Arsenate	276. 25167-82-2	
	162, 143-50-0	Kepone	220. 10124-50-2	Potassium Arsenite	277. 27323-41-7	
	163. 301-04-2	Lead Acetate	221. 7778-50-9	Potassium Bichromate		Dodecylbenzenesulfonate
	164.3687-31-8	Lead Arsenate	222. 7789-00-6	Potassium Chromate	278. 121-44-8	Triethylamine
	165. 7758-95-4	Lead Chloride	223. 7722-64-7	Potassium Permanganate	279. 75-50-3	Trimethylamine
	166. 13814-96-5 167. 7783-46-2	Lead Fluoborate	224. 2312-35-8	Propargite	280. 541-09-3	Uranyl Acetate
	168. 10101-63-0	Lead Fluoride Lead Iodide	225. 79-09-4	Propionic Acid	281. 10102-06-4	Uranyl Nitrate
	169. 18256-98-9	Lead Nitrate	226. 123-62-6	Propionic Anhydride	282, 1314-62-1	Vanadium Pentoxide
	170. 7428-48-0	Lead Stearate	227, 1336-36-3	Polychlorinated Biphenyls	283. 27774-13-6	Vanadyl Sulfate
	171. 15739-80-7	Lead Sulfate	228. 151-50-8 229. 1310-58-3	Potassium Cyanide	284. 108-05-4	Vinyl Acetate
	172. 1314-87-0	Lead Sulfide	230. 75-56-9	Propylana Oxida	285. 75-35-4	Vinylidene Chloride
	173. 592-87-0	Lead Thiocyanate	231. 121-29-9	Propylene Oxide Pyrethrins	286. 1300-71-6	Xylenol
	174. 58-89-9	Lindane	232. 91-22-5	Quinoline	287. 557-34-6	Zinc Acetate
	175. 14307-35-8	Lithium Chromate	233, 108-46-3	Resorcinol	288. 52628-25-8 289. 1332-07-6	Zinc Ammonium Chloride
	176. 121-75-5	Malthion	234. 7446-08-4	Selenium Oxide	290. 7699-45-8	Zinc Borate Zinc Bromide
	177. 110-16-7	Maleic Acid	235. 7761-88-8	Silver Nitrate	291.3486-35-9	Zinc Carbonate
3	178. 108-31-6	Maleic Anhydride	236. 7631-89-2	Sodium Arsenate	292. 7646-85-7	Zinc Chloride
392	179. 2032-65-7	Mercaptodimethur	237. 7784-46-5	Sodium Arsenite	293.557-21-1	Zinc Cyanide
13	180. 592-04-1	Mercuric Cyanide	238, 10588-01-9	Sodium Bichromate	294. 7783-49-3	Zinc Fluoride
	181. 10045-94-0	Mercuric Nitrate	239. 1333-83-1	Sodium Bifluoride	295.557-41-5	Zinc Formate
3	182. 7783-35-9	Mercuric Sulfate	240. 7631-90-5	Sodium Bisulfite	296. 7779-86-4	Zinc Hydrosulfite
	183. 592-85-8	Mercuric Thiocyanate	241. 7775-11-3	Sodium Chromate	297. 7779-88-6	Zinc Nitrate
	184. 10415-75-5	Mercurous Nitrate	242. 143-33-9	Sodium Cyanide	298. 127-82-2	Zinc Phenolsulfonate
	85. 72-43-5	Methoxychlor	243. 25155-30-0	Sodium Dodecylbenzene	299. 1314-84-7	Zinc Phosphide
	86. 74-93-1	Methyl Mercaptan		Sulfonate	300. 16871-71-9	Zinc Silicofluoride
	87. 80-62-6	Methyl Methacrylate	244. 7681-49-4	Sodium Fluoride	301. 7733-02-0	Zinc Sulfate
	88. 298-00-0	Methyl Parathion	245. 16721-80-5	Sodium Hydrosulfide	302. 13746-89-9	Zirconium Nitrate
	89. 7786-34-7	Mevinphos	246. 1310-73-2	Sodium Hydroxide	303. 16923-95-8	Zirconium Potassium Fluoride
	90. 315-18-4	Mexacarbate	247. 7681-52-9	Sodium Hypochlorite	304. 14644-61-2	Zirconium Sulfate
	91. 75-04-7	Monoethylamine	248. 124-41-4	Sodium Methylate	305. 10026-11-6	Zirconium Tetrachloride
		· valuation				n (*)
		64-895-62			2	
	26	2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2				
	*	· 30 - 10 163				

FIELD TRIP SUMMARY REPORT

ORIGINAL (Red)

This summary should be prepared in conjunction with the Preliminary Assessment, EPA Form 2070-12.

EPA Case Number PA - 2087

Site Name U.S. Aluminum Lagoons

Site Description

Facility is located at the southwest corner of Marietta Bouough next to the north bank ofthe Susquehanna River. Facility is a secondary Aluminum Smelting operation; melting scrap Aluminum. Prior to 1975 air emissions were handled by a wet scrubber and waste from from this operation was sent to public sewer. Waste from slag washing operation was sent to two unlined dewatering impoundments. In 1975, U.S. Reduction Co. bought U.S. Aluminum. U.S Reduction replaced the wet scrubber with baghouse and also took the two surface impoundments out of service.

Area of site (acres)
Approx. 5 acres

Hazardous portion, if not entire site agoons = 0.25 acre

Description of processes/operations which took place at the site

Facility melted scrap aluminum in furnace. Prior to 1975 air emissions were handled by a wet scrubber; was replaced by a bag house in 1975. Slag from the smelting operation washed with water to remove soluble components; insoluble material was put back into furnace. This recovery operation was replaced with a dry operation at another plant; the slag is shipped to other plant for recovery. The slag washing operation utilized two earthen dewatering impoundments at southeast corner of the property.

Waste handling/disposal practices

Waste from the wet scrubber was sent to the public sewer. Waste from the bag house is sent off site as a hazardous waste. Waste from the slag washing operation was sent to unlined dewatering impoundments; water and soluble components percolated into the surrounding soil; insoluble components were dredged out periodically and disposed of at undetermined location.

Site topography and runoff drainage pathways

Surface water drains onto the site from the north. The immediate vicinity of the site is relatively flat with surface water draining south towards Susquehanna River.

Surface or subsurface drainage areas (leachate) noted?

Odors/stains noted?

No.

Stressed vegetation noted?

Location and description of streams or receiving waters adjacent to site. Include flow direction and observations. Note location on attached map.

Susquehanna River is approx. 100 yd south ofthe site. The River flows in a west to east direction by the site.

Monitoring wells on site or in vicinity. Note location on attached map.

No.

Population within ¼ mile of site: □ 0-10 □ 10-100 □ greater than 100	Population within 1 mile of site: 0-10 10-100 100-1000 Separate than 1000 (Red)
Surrounding land use (woodlot, agricultura	al, recreation, industrial, etc.)
NORTH Residential	EAST Residential
SOUTH Floodplain; River	WEST Residential Military installation
Two municipal water supply surface intakes Columbia Water Company has surface intake a Water Authority has surface intake just sout	t 2nd St. in Columbia, City of Lancaster
Reference:	
Domestic wells. Approximate number within List nearest wells below and show location	
Owner/Resident Add	dress Phone
Groundwater flow direction, if known	
Description of odor/taste problems	
· · · · · · · · · · · · · · · · · · ·	
State inspection activity (including perm	its held)
DER Bureau of Water Quality involved in clo	sure of the surface impoundments in 1970's.
State/Federal/Private remedial activities	
Impoundments closed by U.S. Reduction and the loading dock.	are now covered over with concrete by

Additional comments--Further description of site

According to company records (see sections 9,10,11,12) the waste that were put into the surface impoundments were not hazardous as defined by RCRA.

The Borough of Marietta has received complaint from nearby residences that air emissions from this facility are corroding the paint off their houses. The waste collected in the baghouse has been found to contain Cd and Pb.

Thie low ranking was given to this site for the possible air emissions and possible soil & surface water contamination by these emissions (Cd & Pb). An assessment of none would have been given just for the surface impuondments due to the waste being non-hazardous as defined by RCRA; although, contamination of groundwater with residual waste is probable.

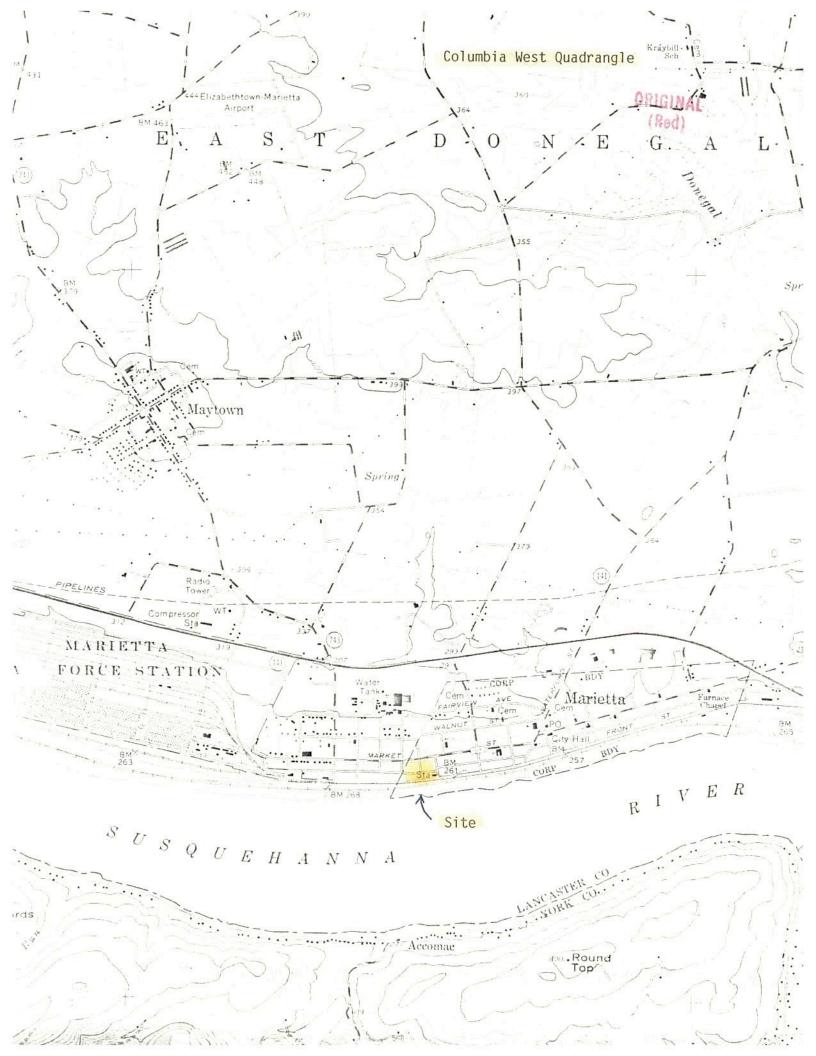
Name and Title	Affili	iation	Phone
Thomas Hendon, Dir. Env./Manufa	cturing Serv.	US Reduction	(800)323-8760
Al Reinhart, Plant Manager	U.S. A1CO		(717)426-1981

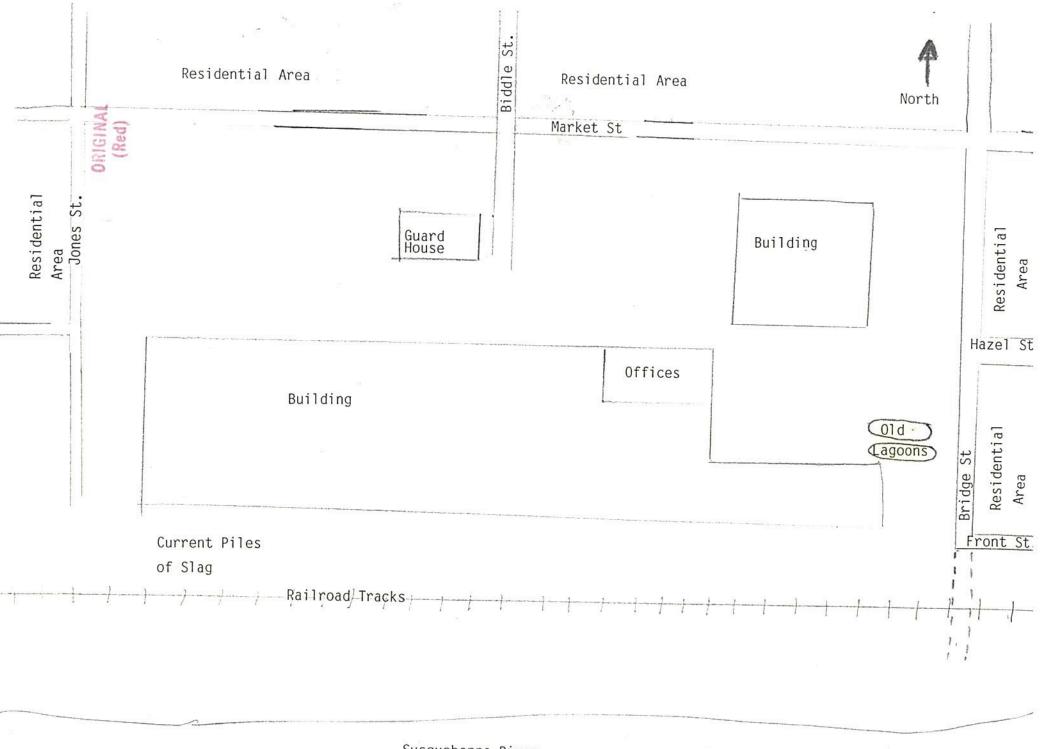
INSPECTION INFORMATION	
Name and title of inspector(s) Thomas Sheehan,	Project Officer
AgencyPa. DER, Div. Emergency & Remedial Response	Phone number <u>(215)565-1687</u>
Date3/19/87	Time on site _5 hour
Weather co.:ditions: Clear, sunny, seasonal temperature.	

ATTACHMENTS

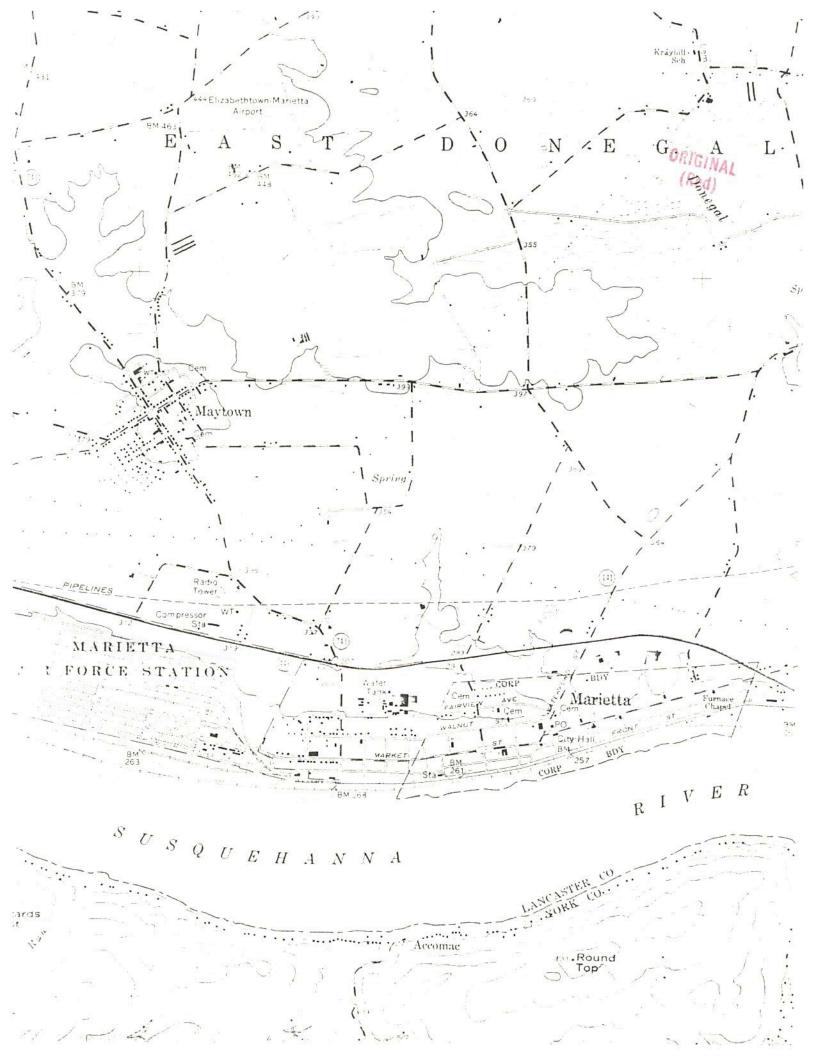
- o Topographic map identifying site location. Include name of quadrangle map. ~
- o Site sketch map showing location of monitoring wells, domestic wells, municipal water supplies, and areas of concern (lagoons, leachate seeps, drums, etc.)
- o Any available sampling results or state monitoring data with map showing sample locations.

fe:





Area Serviced by Matietta Gravity Water Company



Sources of Information:

Commonwealth of Pennsylvania
Department of Environmental Resources
Region 3 Files
One Ararat Blvd.
Harrisburg, Pa. 17110

ORIGINAL (Red)

Recorder of Deeds Lancaster County Courthouse Duke & Orange Sts. Lancaster, Pa. 17603

Lancaster County Planning Commission Lancaster County Courthouse Duke & Orange Sts. Lancaster, Pa. 17603

On site visit 3/19/87 By: Thomas Sheehan, Project Officer Commonwealth of Pa. Department of Environmental Resources Division of Emergency & Remedial Response, Bureau of Waste Management

U.S. ALCO Hazel & Biddle Sts. Marietta, Pa. 17547

Marietta Borough Market Street Marietta, Pa. 17547 (717) 426- 4143

Marietta Gravity Water Company Donnegal Insurance Building Marietta, Pa. 17547 (717)429-1931

USGS 7.5 Minute Quadrangle Topographic

Columbia West Quadrangle . Rev. 1972

Lancaster City Water Authority PO Box 1599 Duke Street Lancaster, Pa. 17603

Columbia Water Company 220 Locust Street Columbia, Pa.

Sources of Information - People Interviewed:

Nancy Parker, Senior Environmental Planner Lancaster County Planning Commission Lancaster County Courthouse Duke and Orange Sts. Lancaster, Pa. 17603

ORIGINAL (Red)

Al Reinhart, Plant Manager U.S. ALCO (Division of U.S. Reduction) Hazel and Biddle Streets Marietta, Pa. 17547 (717)426-1981

Thomas Hendon Corporate Manager of Environmental Response U.S. Reduction Munster, Indiana 1-800-323-8760

William Budding Administrative Assistant Council Marietta Bouough Market Street Marietta, Pa. 17547 (717)426-1981

Miriam Lenhart. Superintendant Marietta Gravity Water Company Donnegal INsurance Building Matietta, Pa. 17547 (717)429-1931

10.



COMMONWEALTH OF PENNSYLVANIA DEPARTMENT OF ENVIRONMENTAL RESOURCES

BUREAU OF WASTE MANAGEMENT One Ararat Boulevard Harrisburg, Pennsylvania 17110 (717) 657-4588 December 10, 1985

ORIGINAL (Red)

Mr. Al Reinhart Plant Manager U.S. Aluminum Corporation P.O. Box 8 Harletta, PA 17547-0008

Re: Wivell Quarry
East Donegal Township
Lancaster County

Dear Mr. Reinhart:

During an investigation of past waste disposal activities at the Wivell Quarry, evidence was uncovered indicating that an industrial waste stream from U.S. Aluminum was deposited here. The following information is requested in order to complete a preliminary assessment of this site.

- 1. Records of the quantities of waste and dates of deposition.
- 2. Specific source of the waste and analyses or expected constituents.
- 3. Identity of the waste transporter, if not U.S. Aluminum.
- 4. Description of the waste handling or disposal practices employed at the quarry.
- 5. Copies of any contracts or agreements with the quarry's owner.

Please supply this information by December 20, 1985.

There are several sources of the statutory authority for requesting this information. The Resource Conservation and Recovery Act, 42 U.S.C. Section 6933, authorizes Pennsylvania to obtain information concerning the amount, nature, and toxicity of the hazardous substances which may have been stored or disposed of at any time. The Comprehensive Environmental Response Compensation

12 Rood records destroyed

Strye ava - stow Runch.

. 2 of 2

ORIGINAL (Red)

Mr. Al Reinhart

-2-

December 10, 1985

and Liability Act ("Superfund") also authorizes the Department to obtain this information. Sections 103 and 104 of the Superfund Act, 42 U.S.C. Sections 9603 and 9604 authorize the state to obtain and require persons who have stored or disposed of hazardous substances to furnish information relating to such substances as specified in those sections. Finally, the Pennsylvania Solid Waste Management Act, 35 P.S. Section 6018.104, 6018.502(f) and 6018.603 authorizes the Department to obtain information concerning storage and disposal of solld waste.

If you have any questions concerning this matter, please feel free to contact me.

Sincerely,

Gregory L. Harder

Solid Waste Specialist Harrisburg Regional Office

GLH: flw

CC: Thomas Sheekan 4/21/87

DEPARTMENT OF ENVIRONMENTAL RESOURCES

DEPARTMENT OF U.S. REDUCTION CO.

2025 APR 2 4 1987 LANSING, ILLINOIS 50438

(312) 895-9400 TELEX: 206669 ALUMINUM ALLOYS FOR EVERY PURPOSE (Red)

DELAWARE COUNTY OFFICE

January 2, 1986

Mr. Gregory L. Harder Commonwealth of Pennsylvania Department of Environmental Resources Bureau of Waste Hanagement One Ararat Boulevard Harrisburg, PA 17110

Dear Mr. Harder:

This letter is in response to your letter to Mr. Al Reinhart dated December 10, 1985 and our telephone conversation of December 16, 1985 regarding the disposal activities at the Wivell Quarry. We appreciated the extension to January 8, 1986 to respond to your information request.

It is our understanding that the grab sample taken from the Wivell Quarry contained high quantities of total metals(i.e. cadmium, lead, copper, aluminum, magnesium, chromium, iron, manganese, etc...). Also, when the sample was run for E. P. toxicity the only metal that would constitute a potential hazard in accordance with R.C.R.A. was selenium.

You mentioned that there were records showing U. S. Aluminum Corp. disposed of dust in the Wivell Quarry in 1976. The Company has not found any records to indicate that it made waste deposits at the Wivell Quarry.

After the acquisition of U. S. Aluminum Corp. by U. S. Reduction Co., an agreement with the Pennsylvania D.E.R. was entered into to shutdown the wet mill operation along with removal of the material which had accumulated on the company's property from this operation. This material consisted of a dust pile. Presently there are no records to ascertain the disposition of this pile.

If it is assumed for the sake of discussion that some of this dust was disposed of at Wivell Quarry, the information in answer to you questions is as follows:

 There are no records of the quantity of waste and dates of deposition. Wivell Quarry Page 2 January 2, 1986



- 2. There is no analysis of this material per se(see analysis discussion at end of letter).
- Identity of the waste transporter is not known.
- 4. We do not know about the waste handling or disposal practices employed at the Quarry.
- After a diligent search, we have been unable to find copies of any contract or agreement with the Quarry's owner.

I would like to offer the following background information on the wet mill operation along with analyses of the slags in our industry, and work done with the Aluminum Recycling Association on Secondary Aluminum High Salt Slag which resulted in removing from the R.C.R.A. Hazardous Listing.

WET MILL OPERATION:

O.H. Slag known also as High Salt Slag and sometimes Aluminum Dross Slag is a basic mixture of Aluminum Oxide 40-55%, Sodium Chloride, and Potassium Chloride at 35-40%, and Aluminum Metal at 10-15%. The purpose of the wet mill is to concentrate the aluminum metal so it can be recycled to the furnace for recovery. The wet mill does this by washing/disolving the soluable sodium and potassium salts freeing up the aluminum and breaking by tumbling and reducing the insoluable oxides to a smaller size fraction that can be carried away with the wash water. The oversize concentrate is screened out and dried. The water effluent with the dissolved salts and suspended solids were discharged to a settling lagoon, where the soluable salts as total dissolved solids percolated through the ground underlying the lagoons. As the lagoons filled up with undissolved suspended solids, they were dredged out. The residue dust was piled until disposed of.

It would be apparent that the brine leaching process would have leached out most of the leachables leaving inert aluminum oxide residue containing a $\underline{\text{minor}}$ concentration of aluminum metal approximating 1-3% by weight.

In my telephone conversation with you, you indicated a R.C.R.A. E.P. toxicity was run on the Quarry sample and the only toxic substance according to R.C.R. A. was Selenium that exceeded the limitations. In all of the R.C.R.A. E.P. toxicity leach tests we've conducted on our O.H. slag, selenium has never been found in any significant concentration.

As you know, there is a trace amount of selenium in our furnace baghouse dust. This is most likely due to concentrating by vaporization.

at the melting furnace operating at 1450° F and condensing back to a particulate captured by the baghouse. These baghouse dusts analyzed for E.P. toxic metals have never exceeded RCRA for selenium.

The following analytical information is itemized below and enclosed for your records on U. S. Aluminum Corp:

- 1. Analysis of Secondary Aluminum Smelting Waste Products for U. S. Reduction Co. by Van Note-Harvey 1979(Excerpts of Corporate Report). NOTE: The RCRA proposed limitations during this report is now ten(10) times greater as the promulgated standard.
- 2. I.U. Conversion study analyses(Excerpts from "Waste Management Assessment for U. S. Reduction Co. Marietta, Pennsylvania 1979). NOTE: The RCRA Standard limitations are ten(10) times greater. Also note that Mix 2 and Mix 8 would have included baghouse dust and other I.U. Conversions waste and/or additives.
- 3. Aluminum Recycling Associations work and report got the Federal EPA to delist the Secondary Aluminum Industry "High Salt Slag" from the Hazardous Waste Listing.

A copy of this report is enclosed. Note also that this work was done before the final regulations were promulgated, therefore, the limitations then should be multiplied by ten(10) to yield the present standard today.

I hope this information will help you in your preliminary assessment of the Wivell Quarry. If I can be of any further help, please don't hesitate to contact me at (312) 895-9400 ext. 283.

Respectfully,

U. S. REDUCTION CO.

Thomas R. Hendon, P.E.

Director Environmental/Manufacturing Services

TRH/jah Enclosures

ITEM 1

ORIGINAL (Red)

ANALYSIS OF SECONDARY ALUMINUM
SMELTING WASTE PRODUCTS
FOR
U. S. REDUCTION COMPANY

Prepared for
U. S. Reduction Company
East Chicago, Indiana 46312

Prepared by
Van Note-Harvey Associates
Consulting Engineers and
Environmental Laboratories

1531 North Main Street Blacksburg, Virginia 24000

September 20, 1979



TABLE V-1

U. S. Reduction Slag and Dust Waste Samples

LOCATION	TYPE WASTE MATERIAL	PRODUCTION DATA
Marietta. Pennsylvania	High salt slag	Six ladles taken of hot slag from Furnaces 1, 2, and 3 at USALCO. Represents 2,429 lbs. slag. Sent 6/15/7
Fontana. California	High salt slag	Ten ladles of hot slag taken from Furnaces 51 and 52. Represents 12,460 lbs. slag. Sent 6/15/70
Toledo, Chio	High salt slag	Thirty-five ladles of hot slag taken from 7 alloys on Furnaces 23, 25 and 26. Represents 10,500 lbs. slag. Sent $6/18/79$
Alton. Illinois	High salt slag	Twelve ladles of hot slag taken from Furnaces 2, 7 and 3. Represents 12 slag pans or 18,000 lbs. slag. Sent 0/15/79
Alton, Illinois	Mesh dust pile	Composite cross section sample taken by J. Gordon. Sent $7/13/79$
East Chicago, Indiana	High salt slag	Ten ladles of hot slag taken from Furnaces 1, 2, and 3, and broken up hot to split on $\frac{1}{2}$ " riffle. Represents 17,500 lbs. slag. Sent $6/13/79$
East Chicago, Indiana	Mesh dust, East Chicago plant	Representative composite sample from Lot 380. Represents 100,455 lbs. of slag. Sample weight 2108 grams. Sent 6/14/79
East Chicago, Indiana	Mesh dust, Reynolds-McCook	Representative composite sample of 3 truck lots from primary supplier. Represents 115,390 lbs. of slag. Sample weight 400 grams. Sent 0/14/79
Russellville, Alabama	High salt slag	Twelve ladles taken from Furnaces 36, 37, and 38. Represents 18,000 lbs. of slag. Sent 6/15.79
Russellville, Alabama	Mesh dust, milled 0/14/79	Representative composite sample from Lot 0589 sampled from tote boxes on $6/14/79$. Represents $73,739$ lbs. of slag.
Russellville, Alabama	Mesh dust, Main Old Pile	Composite sample taken the week of 7 9/70 from old leached dust pile.
Russellville. Alabama	Mesh dust, New Pile	Composite sample taken the week of 7 0/70 from currendust pile.



TABLE V-2

Comparison of Four Extractions of Alton Mesh Dust Pile

Extract Levels (mg/l) Mean and				Mean and Standard	
Contaminant	Extract #1	Extract =2	Extract #3	Extract #4	Deviation
Arsenic	0.001	0.002	<0.001	<0.001	NA
Barium	3.4	0.50	0.70	0.52	1.28 = 1.42
Cadmium	<0.02	<0.02	<0.02	<0.02	NA
Çhromium	0.10	0.10	0.24	0.14	0.10 = 0.00
Lead	0.30	0.25	0.32	0.24	0.28 = 0.04
Mercury	0.001	0.001	<0.001	<0.001	NA
Selenium	<0.001	<0.001	<0.001	<0.001	NA
Silver	<0.01	<0.01	<0.04	<0.04	NA
Zinc	7.5	0.0	11.2	7.1	8.10 = 2.1
Copper	6.8	6.4	10.3	6.4	7.6 = 2.1
Nickel	0.10	0.20	0.40	0.28	0.32 = 0.10
Sodium	53	58	14	10	34 - 25
Potassium	12	15	26	21	10 = 6
Magnesium	410	160	290	160	255 = 120
Chloride	70	50	110	So	78 = 25
Amount of Acetic Acid Required to Stabilize pH (ml)	238	159	1 54	1 57	

Note: NA - Not applicable since levels are below detectable limit.

TABLE V-3

Comparison of Three Digested Aliquots of Original Extract of Alton Mesh Dust Pile

ORIGINAL (Red)

Contaminant	Ext. Digestion =1	ract Levels (m	—	Mean and Standard
	DIECECTOR =1	Digestion #2	Digestion =3	Deviation
Arsenic	<0.001	< 0.001	<0.001	NA
Barium	3.4	0.9	0.9	1.7 = 1.4
Cadmium	<0.02	<0.02	<0.02	NA
Chromium	0.16	0.15	0.13	0.15 = 0.02
Lead	1.20	0.30	0.40	0.0 = 0.5
Selenium	<0.001	<0.001	< 0.001	NA
Silver	<0.04	<0.01	<0.01	NA
Zinc	7.5	7-4	7.2	7.4 = 0.15
Copper	ó.S	7.1	6.7	6.0 = 0.2
Nickel	0.10	0.40	0.45	0.42 = 0.03
Sodium	53	11	50	40 = 5
Potassium	12	14	15	
Magnesium	410	290	250	14 = 1.5 320 = 80
				~

3

NA - Not applicable since levels are below detectable limit.

TABLE V-4

Extraction Procedure Results

Control Sample Test of Leaching from Extraction Equipment

Contaminant	Extract Level, mg/l
Arsenic	<0.001
Barium	<0.25
Caimium	<0.02
Lead	<0.05
Mercury	<0.001
Selenium	<0.001
Silver	<0.04
Zinc	<0.01
Copper	<0.05
Nickel	<0.05

TABLE V-6

Extraction Procedure Results

Marietta, Pennsylvania High Salt Slag

Contaminant	Extract Level	Allowable Extract Level Under EPA Proposed Regulations. mg/l X/O
Arsenic	0.002	0.50
Barium	2.6	10.0
Cadmium	< 0.02	0.10
Chromium	< 0.05	0.50
Lead	0.06	0.50
Mercury	< 0.001	0.02
Selenium	< 0.001	0.10
Silver	< 0.01	0.50
Zinc	< 0.1	
Copper	< 0.05	
Nickel	< 0.05	
Sodium	4830	
Potassium	4410	
Magnesium	530	
Chloride	9540	

TABLE V-7

Extraction Procedure Results

Fontana, California High Salt Slag

Contaminan	Extract =1	ract Levels (Extract =2	mg (1) Extract =3	Allowable Extract Level Under EPA Proposed Regulations. mg/l	XIO
Arsenic	<0.001	0.001	<0.001	0.50	STALL SI
Barium	3.3	5.5	×\$	10.0	
Cadmium	<0.02	<0.02	<0.02	0.10	
* Chromium	<0.05	43-	**	0.50	
Lead	<0.05	(49)	A.	0.50	
Mercury	0.028	<0.001	<0.001	0.02	
Selenium	<0.001	<0.001	<0.001	0.10	
Silver	<0.01	<0.01	<0.01	0.50	
line	0.7	0.2	0.2		
Copper	<0.05	<0.05	<0.05	-	
Nickel	<0.05	<0.05	<0.05		
Sodium	5600	**	**		
Potassium	3730	43	24		
Magnesium	990	÷.	2.5	50-60 ⁴ (2000)	
Chloride	10,300	ist.	**	50.55 0	

Note: --- No limit currently specified by EPA. Parameter not rechecked.

TABLE V-8

Extraction Procedure Results

ORIGINAL (Red)

Toledo, Ohio High Salt Slag

Contaminant	Extract Level	Allowable Extract Level Under EPA Proposed Regulations. mg/l X/O
Arsenic	0.003	0.50
Barium	5.0	10.0
Cadmium	<0.02	0.10
Chromium	<0.05	0.50
Lead	0.10-	0.50
Mercury	<0.201	0.02
Selenium	<0.001	0.10
Silver	<0.01	0.50
Zinc	1.0	
Copper	0.10	
Nickel	0.15	
Sodium	5390	
Potassium	3010	
Magnesium	340	
Chloride	9500	

Note: ---No limit currently specified by EPA.

Average value of two separate extracts.

TABLE V=0

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ORIGINAL (Red)

Extraction Procedure Results

Alton, Illinois High Salt Slag

Contaminant	Extract Level	Allowable Extract Level Under EPA Proposed Regulations, mg/1 X 10
Arsenic	ক.001	0.50
Barium	6.7	10.0
Cadmium	<0.02	0.10
Chromium	0.03	0.50
Lead	0.25	0.50
Mercury	a.001	0.02
Selenium	<0.001	0.10
Silver	<0.01	0.50
Zinc	5.3	
Copper	0.07	9 7
Nickel	0.10	
Sodium	6800	
Potassium	4820	
Magnesium	210	
Chloride	10,000	



TABLE V-10

Extraction Procedure Results

Alton, Illinois

Mesh Dust Pile

Contaminant	Extract Level	Allowable Extract Level Under EPA Proposed Regulations. mg/l X O
Arsenic	<0.001	0.50
Barium	3.4	10.0
Cadmium	<0.02	0.10
Chromium	0.10	0.50
Lead	0.30	0.50
Mercury	0.001	0.02
Selenium	<0.001	0.10
Silver	<0.01	0.50
Zinc	7.5	
Copper	6.3	
Nickel	0.4	
Sodium	53	
Potassium	12	
Magnesium	410	
Chloride	70	
Sulfate	136	
Total Dissolved Solids	2510	

Note: ---No limit currently specified by EPA. "Average of four separate extracts.

TABLE V-11

Extraction Procedure Results



East Chicago, Indiana High Salt Slag

Contaminant	Extract Level	Allowable Extract Level Under EPA Proposed Regulations. mg/l X 10
Arsenic	< 0.001	0.50
Barium	1.8	10.0
Cadmium	< 0.02	0.10
Chromium	0.05	0.50
Leai	0.2	0.50
Mercury	< 0.001	0.02
Selenium	< 0.001	0.10
Silver	< 0.01	0.50
Zinc	10.0	
Copper	0.5	
Nickel	0.2	
Sodium	4710	===
Potassium	3440	
Magnesium	300	
Chloride	7100	

TABLE V-12

(Red)

Extraction Procedure Results

East Chicago, Indiana Mesh Dust

-East Chicago Plant-

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Contaminant	Extract Level mg/l.	Allowable Extract Level Under EPA Proposed Regulations. mg/l X O
Arsenic	0.001	0.50
Barium	7.3	10.0
Cadmium	< 0.02	0.10
Chromium	<0.05	0.50
Lead	<0.05	0.50
Mercury	0.002	0.02
Selenium	<0.001	0.10
Silver	<0.01	0.50
Zinc	3.2	
Copper	0.15	
Nickel	0.10	
Sodium	4650	
Potassium	4620	
Magnesium	230	
Chloride	10.000	

TABLE V-13

(Red)

Extraction Procedure Results

East Chicago, Indiana Mesh Dust

-Reynolds-McCook-

Contaminant	Extract Level	Allowable Extract Level Under EPA Proposed Regulations. mg/l XIO
Arsenic	< 0.001	0.50
Barium	0.7	10.0
Cadmium	< 0.02	0.10
Chromium	< 0.05	0.50
Lead	< 0.05	0.50
Mercury	< 0.001	0.02
Selenium	< 0.001	0.10
Silver	< 0.04	0.50
Zinc -	11.6	
Copper	0.38	
Nickel	< 0.05	
Sodium	315	
Potassium	482	
Magnesium	750	
Cnloride	1520	

TABLE V-14

Extraction Procedure Results

Russellville, Alabama High Salt Slag

Contaminant	Extract Level mg/l	Allowable Extract Level Under EPA Proposed Regulations, mg/l X/C			
Arsenic	< 0.001	0.50			
Barium	2.2	10.0			
Cadmium	< 0.02	0.10			
Chromium	0.15	0.50			
Lead	0.15	0.50			
Mercury	0.002	0.02			
Selenium	< 0.001	0.10			
Silver	< 0.01	0.50			
Zinc	1.7	1			
Copper	0.05				
Nickel	0.05				
Sodium	5700				
Potassium	1000				
Magnesium	110	()			
Chloride	10,000				

TABLE V-15
Extraction Procedure Results

ORIGINAL (Red)

Russellville, Alabama Mesh Dust -MILLED 6/14/70-

Contaminant	Extract Level mg/l	Allowable Extract Level Under EPA Proposed Regulations. mg/l X O
Arsenic	< 0.00.1	0.50
Barium	3.3	10.0
Cadmium	< 0.02	0.10
Chromium	< 0.05	0.50
Lead	0.25	0.50
Mercury	< 0.001	0.02
Selenium	< 0.001	0.10
Silver	< 0.01	0.50
Zinc	2.3	
Copper	0.27	
Nickel	0.12	
Sodium	4260	
Potassium	4110	
Magnesium	100	
Chloride	3600	

TABLE V-10

Extraction Procedure Results

Russellville, Alabama Mesh Dust

-Main Old Pile-

	Extract Level	Allowable Extract Level Under EPA Proposed
Contaminant	mg/l	Regulations. mg/l X/C
Arsenic	0.002	0.50
Barium	1.1	10.0
Cadmium	0.056	0.10
Chromium	< 0.05	0.50
Lead	0.10	0.50
Mercury	0.003	0.02
Selenium	< 0.001	0.10
Silver	< 0.01	0.50
Zinc	2.2	
Copper	4.1	
Nickel	0.02	
Sedium	320	
Potassium	223	
Magnesium	240	
Chloride	430	

TABLE V-17 Extraction Procedure Results



Russellville, Alabama Mesh Dust

-New Pile-

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Extract Level mg/l	Allowable Extract Level Under EPA Proposed Regulations, mg/l X 10
< 0.001	0.50
1.7	10.0
< 0.02	0.10
< 0.05	0.50
0.2	0.50
0.001	0.32
< 0.001	0.10
< 0.01	0.50
4.2	
9.1	
0.28	
483	===
195	
135	
700	
	mg/1 <0.001 1.7 <0.02 <0.05 0.2 0.001 <0.001 <0.04 4.2 9.1 0.28 483 195 135

Section 11

2

ITEM 2

FOR
U.S. REDUCTION COMPANY
MARIETTA, PENNSYLVANIA

N. P.

IU Conversion Systems, Inc. 115 Gibraltar Road Horsham, Pa. 19044

TABLE V

ASTM (A) LEACHATE ANALYSIS

ORIGINAL (Red)

Param	eter	Baghouse Dust	MgCl ₂ Slag	FCE(OH) Slag	Mix 2	Mix 8		RCRA Standard X 10
рН		4.2	10.1	9.9	8.5	6.89	ři,	-
TDS		108,476	28,648	162,197	15,030	96,700		55 53 0
Ag		.25	<.05	.05	<.05	.14		0.50
As		44	.02	.02	.01	.21		0.50
Ва		<10	<.10	< :10	< .10	.70	d	10.0
ca,		45.0*	<.01	<.01	<.01	5.40*		0.10
Cr		0.23	<.05	<.05	< .05	.10		0.50
Cu		-	.,,=	-	_	.25		-
Fe .		· -		_	:=	.69		
Ηg		.0020	<.001	.0 < .0010	<.0010	.001		0.02
Mn		-	()	-		3.07	5	-
РЪ		17.6*	<.05	<.05	<.05	1.28*		0.50
Se		<.01	<.01	.01	.13*	<.05		0.10
Zn		-	_	-	-	1.26		-

Results reported in ppm except pH

14

^{*} Exceeds RCRA Standard for Toxic Waste

Section 12

ITEM 3

Statement of the Aluminum Recycling Association In Response to a Notice of Proposed Rule-Making Under Section 3001 of the Solid Waste Disposal Act, As Amended

Recovery Act of 1976.

Submitted to the Office of Solid Waste
U.S. Environmental Protection Agency
Washington, D.C. 20460

by the Resource Conservation and

March 16, 1979

Richard M. Cooperman

Executive Director
Aluminum Recycling Association
900 - 17th Street, N.W.
Washington, D.C. 20006
(202) 785-0550

ORIGINAL

The Aluminum Recycling Association represents most ofed) the production capacity of the aluminum recycling firms in this country. Member companies range from divisions of some of the nation's largest companies to medium-sized and small, family-owned businesses. Over 5,000 people are employed in the nearly 45 industrial plants operated by these firms.

One thing in common to these member firms is their commitment to maximizing the useful life of the world's steadily diminishing mineral reserves through the recycling of aluminum. As such, the values of the industry are fully consonant with the goals of the Resource Conservation and Recovery Act: mineral recovery through recycling, environmental protection through conservation of virgin mineral sources, and energy conservation and economic benefits through re-use of existing sources.

Section 1002 of the Act, 42 U.S.C. 6901, provides in part:

"(c) Materials. -- The congress finds with respect to materials, that --

"(1) millions of tons of recoverable material which could be used are need-lessly buried each year;

"(2) methods are available to separate usable materials from solid waste; and

"(3) the recovery and conservation of such materials can reduce the dependence of the United States on foreign resources and reduce the deficit in its balance of payments.

"(d) Energy. -- The Congress finds with respect to energy, that --

- "(1) solid waste represents a potential source of solid fuel, oil, or gas that can be converted into energy;
- "(2) the need exists to develop alternative energy sources for public and private consumption in order to reduce our dependence on such sources as petroleum products, natural gas, nuclear and hydroelectric generation; and
- "(3) technology exists to produce usable energy from solid waste."

Today, aluminum recycling is one of the pre-eminent environmental industries in the United States. Approximately 75 recycling plants throughout the country re-process aluminum scrap - from industrial waste to used cars - into clean and re-usable metal. Thus, the industry converts what would otherwise constitute

a solid waste disposal problem of great magnitude for communities across the land, not to mention an ecological eyesore of gross proportions, into economically, socially and environmentally desirable use. The success of the industry, measured by its annual sales between 700 and 800 million dollars, exposes the myth that the economy and the environment are mutually exclusive values.

Most of the recycling plants in operation today utilize only high-grade aluminum scrap with little attendant waste. However, between 25 and 35% of total secondary aluminum produced in this country each year involves the recovery of low-grade aluminum scrap through a process known as dross smelting. Recycling of the low-grade scrap, which generally contains between 10 and 30% aluminum, requires a two-step smelting process to maximize the metal recovery. The original smelting, which is performed in reverbatory furnaces, recovers approximately 75% of the aluminum content in the scrap; the dross waste is itself later recycled in a second process which increases the metal recovery to approximately 96-97%. A few plants, which generally utilize the lowest grade aluminum scrap, use rotary furnaces; the rotary process is similar to the reverbatory, except that generally only one recycling is necessary to produce between 95-96% metal recovery. The rotary process also produces a dross slag concentrated in larger-sized aggregate chunks than the more traditional reverbatory process. In both processes, fluxing agents and alloying agents are charged with the scrap and later recovered by dissolution in water or as a flux salt. Much of this is later recycled as a steel melting flux cover agent, while other additional material is sold to the cement industry. The major source of solid waste arises from both the impure constituents in the enriched dross and the salt flux used in the smelting process. Arguably the aluminum dross smelting process is among the most environmentally laudable, in that it results in the valuable mineral recovery of what clearly would otherwise line our waste land-fills, not to mention our roadsides.

The environmental pre-eminence of the industry, however, is threatened by the recent proposed rules issued by EPA pursuant to Sections 3001, 3002 and 3004 of the Solid Waste Disposal Act, as amended by the Resource Conservation and Recovery Act. These proposed rules, as explained below, incorrectly classify secondary aluminum dross smelting as an industrial process which results in the generation of hazardous waste (high salt slag plant residue). As a listed "hazardous waste", the proposed rules, if approved, will subject the industry to prohibitively expensive regulation such as to result in significant recycling plant closures and, thereby, a net decrease in overall environmental benefit.

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"hazardous waste" as:

A solid waste, or combination of solid wastes, which because of its quantity, concentration, or physical, chemical or infectious characteristics may--

- (a) cause or significantly contribute to an increase in mortality or an increase in serious irreversible, or incapacitating reversible, illness; or
- (b) pose a substantial present or potential hazard to human health or the environment when improperly treated, stored, transported, or disposed of, or otherwise managed.

In order to provide guidance to producers as to whether their waste is hazardous, Section 3001(a) of the Act, 42 U.S.C. 6921, authorizes the Administrator of EPA to promulgate rules to identify the criteria for identifying the characteristics of and for listing hazardous waste. Section 3001(b) then provides two mechanisms for determining whether a waste is hazardous: a set of characteristics of waste and a list of particular hazardous wastes.

In its proposed rules, EPA has identified several characteristics of hazardous waste (ignitability, corrosivity, toxicity, radioactivity, infectiousness, phytotoxicity, and teratogenicity and mutagenicity) and has set scientific standards respectively. A waste is identified as hazardous either because it exhibits one of the above characteristics at a level above the waste. Both particular wastes and sources or classes of waste streams appear on the hazardous waste list. Section 250.12(b) of of waste be listed if the waste:

(1) possesses any of the characteristics identified in proposed 40 C.F.R. §250.13, and/or



extract of which contains certain contaminants at a level equal to or above ten times the EPA National Interim Primary Drinking Water Standards for the respective substances.

I STATE CO.

I. Secondary Aluminum Dross Smelting Does Not Produce Contaminants Above the EPA Standards for Hazardous Waste, Does Not Meet the Statutory Definition of Hazardous Waste, And Therefore Should be Taken Off the List of Hazardous Wastes.

Section 250.14 of the proposed rules provides that a listed hazardous waste shall not be considered hazardous if it can be demonstrated, pursuant to procedures under Section 250.15, that the Administrator's basis for listing the waste does not meet the criteria for listing under Section 250.12(b). The Aluminum Recycling Association recently contracted with Herron Testing Laboratories, Inc. of Cleveland, Ohio, for analysis of a representative sampling of high salt slag plant residue from five aluminum recycling plants utilizing dross smelting reprocessing procedures. The overwhelming weight of evidence from this analysis indicates that the waste generated by secondary aluminum dross smelting does not reach the standards proposed by EPA to establish toxicity. The analysis is appended at the end of this statement.

The data is summarized as follows:

- (1) The proposed rules establish a maximum allowable toxicity for silver as 0.50 mg/l in the waste extract. The level of toxicity for the tested samples ranged from a low of less than 0.01 mg/l to a high of 0.08 mg/l, a maximum of 16% of the allowable standard.
- (2) The proposed rules establish a maximum allowable toxicity for barium as 10.0 mg/l. The level of toxicity for the tested samples ranged from a low of less than 1.0 mg/l to a high of 5.5, a maximum of 55% of the allowable standard.
- (3) The proposed rules establish a toxic standard for cadmium as 0.10 mg/l. The tested samples ranged from a low of 0.03 mg/l to a high of 0.07, a maximum of 73% of the allowable standard.
- (4) The proposed rules establish a toxic standard for arsenic as 0.50 mg/l. The tested samples ranged from a low of less than 0.001 mg/l to a high of 0.004, a maximum of .8% of the allowable standard.

- (5) The proposed rules establish a toxic standard for mercury as 0.02 mg/l. The tested samples ranged from a low of less than 0.0002 mg/l to a high of 0.0007, a maximum of .035% of the allowable standard.
- (6) The proposed rules establish a toxic standard for selenium as 0.10 mg/l. The tested samples uniformly ranged below 0.01 mg/l.
- (7) The proposed rules establish a toxic standard for chromium as 0.50 mg/l. The tested samples ranged from a low of less than 0.02 mg/l to a high of 0.13, a maximum of 26% of the allowable standard.
- (8) The Proposed rules establish as a toxic standard of 0.50 mg/l for lead. Exclusive of one sample of aluminum scrap waste which is discussed below, the tested samples generally meet the toxic standards by wide margins. While 10 test runs of plant sites A-D averaged less than 0.18 mg/l of lead, or less than 36% of the proposed EPA standard, the first sample taken at Plant E (Lab. No. 7266) tested out at levels significantly higher than the EPA proposed standard. Further inquiry disclosed the following pertinent facts: (1) Plant E utilized a rotary furnace, in contradistinction to Plants A-D, as well as to the vast majority of plants throughout the industry, and, for economic reasons, did not re-process the slag dross produced from the initial smelting. (2) Plant E, like most rotary furnace operations, utilized low-grade aluminum scrap, including high lead content non-magnetic fines of auto-shredded scrap.

The above facts are pertinent because they strongly suggest that the initial sample taken at Plant E is clearly an aberration and should not be taken as representative of an entire industry which generally produces waste well within the EPA proposed guidelines. This conclusion is supported by a second sampling taken at Plant E (Lab. No. L8369), eliminating the nonmagnetic auto-shredded scrap. In addition, the new dross was further crushed into -20 mesh, a particle size roughly equivalent to the dross produced in standard reverbatory operations. retested dross was found to be well within the EPA proposed standards, with the two additional test runs leveling out at 0.11 mg/l and 0.08 mg/l of lead, respectively, or an average of less than 20% of the allowable standard. It is suggested that, were rotary furnace operations, such as Plant E, to process a higher grade of aluminum scrap, or in the alternative, to crush its smelted waste to the same particle size as the slag dross produced from reverbatory operations, they too would consistently meet the proposed EPA criteria.

ORIGINAL

Moreover, even EPA's own data does not appear to support listing of the industry as a generator of hazardous waste. The 1977 CALSPAN Study, which served as the basis of EPA's proposed action in this matter, cited the industry as producing high levels of lead and chromium in its waste. However, apparently only one sampling of high salt slag was taken at a single site, which tested out at 0.24 mg/l of lead (less than half of the allowable standard) and 1.5 mg/l of chromium. As discussed above, the Herron Laboratories analysis conducted at five different sites did not disclose a single sample to be above the proposed EPA toxic standard for chromium; in fact, out of 12 separate test runs, the highest concentration of chromium amounted to only 26% of the allowable standard. Clearly, the CALSPAN date, which is taken from a single source and which does not even violate the proposed lead standard, cannot be used to justify regulation of an entire industry.

furnace operations are the distinct minority in the industry, they perform a valuable environmental and waste recovery function. Utilizing low grade scrap which would otherwise go unrecovered, the Plant E-type rotary operations do the "dirty work" in the industry that no one else would touch. They are also the most marginally profitable. As such, they should be encouraged and given special consideration, not driven out of business by prohibitive regulation.

Clearly the overwhelming weight of the evidence, as indicated by the Herron Laboratories test findings, does not support the proposed EPA finding that the entire secondary aluminum dross smelting industry be listed as a generator of hazard-In the face of this additional technical data, it ous waste. would be arbitrary and unreasonable to so conclude. The one lab sample which tested above the proposed standard for lead must simply be viewed as an aberration - insufficient by itself to subject an entire industry to costly regulation; the inequity of the result becomes even more compelling when it is recognized that single step rotary furnace operations which utilize low grade aluminum scrap perform a socially and environmentally desirable function which would otherwise be neglected. Accordingly, the ARA respectfully requests that secondary aluminum dross smelting - high salt slag plant residue be taken off of the list of hazardous wastes in any final rules promulgated.

II. The Burden of Compliance With the Hazardous Waste Regulations Would Disrupt the Cost-Effectiveness of Low-Grade Aluminum Recycling, Force Plant Closures and Lay-Off of Employees, Result in Reduced Recycling of Aluminum Scrap, and Reduce Net Environmental Protection.

In 1978, the secondary aluminum industry produced nearly 2.5 billion pounds, or approximately 1.1 million metric tons, of specification aluminum alloy out of industrial and consumer waste scrap. The industry, which accounts for a quarter of the country's aluminum needs, takes billions of pounds of scrap out of the waste streams and recycles it into ingots for later die casting and foundry uses. Recycled aluminum eventually finds itself in manufactured automobile parts, tools, machinery and other consumer goods.

Vation advantages to recycling aluminum. The per unit costs are substantially less than for the production of primary aluminum. By definition, the recycling industry furthers environmental values, by converting ecological eyesores and other waste problems into clean metal and useful products. Moreover, the industry is fully consonant with our national goals of energy conservation, by utilizing only 5% of the energy per unit of aluminum as required by the primary industry.

The secondary aluminum industry also conserves other important raw materials. To produce one ton of primary aluminum, the following materials are required: 4 tons of bauxite, 1000 pounds of soda, 250 pounds of lime, 1500 pounds of petroleum coke and pitch, 60 pounds of cryolite, 80 pounds of aluminum fluoride, and 14,000 kilowatts of electricity. To produce one ton of secondary aluminum requires none of the above constituent elements and only 700 kilowatts of electricity.

The environmental efficiency of the industry is further enhanced by the nature of the recycling process. Not only is primary waste and scrap processed, but the resulting waste of the initial recycling in the standard reverbatory operation is itself re-processed in an attempt to maximize aluminum recovery.

As in any industrial process, the recycling of aluminum scrap results in waste, which is called sludge or dross. The dross is composed of the constituent elements of the original scrap, plus aluminum residue which could not be recovered in the initial smelting process. The dross produced by reverbatory furnaces is then further processed to increase the aluminum yield. The secondary dross, together with recovered fluxing and alloying agents, combine to produce a high-salt slag dross residue.

(Red)

This resulting waste is then stored in large slag piles, most of which are located on the recycling premises. A small percentage of the dross is sold to the cement industry for further productive use, certain flux salts are themselves recycled into steel smelting flux cover agents, and other chemicals are also recovered for later commercial uses. But while the industry is moving towards zero waste recycling, the complete recovery of materials from the dross is neither economically nor technically feasible at the present time.

The amount of unrecovered dross which remains as waste is significant. According to one CALSPAN Study commissioned by EPA, 1400 Kg. of dross waste is produced for every 1000 Kg. of recycled aluminum. Thus the dross waste disposed of in 1978 alone is estimated at 1.5 million metric tons, with accumulated waste stored at approximately 75 plant sites across the United States conservatively estimated at over 10 million metric tons. For example, one company estimates that its 14 acres of dross contains over 160,000 metric tons at one plant site alone, which increases at a rate of 300 metric tons for each day of plant operation.

The cost of compliance with even part of EPA's proposed hazardous waste regulations for the disposal of such large volumes of high-salt slag dross would be prohibitive and would result in significant plant closures and reduced recycling of secondary aluminum. While most dross piles are sited far away from public drinking water sources, few are lined with clay or lime. The cost of constructing lined land fills to meet EPA specifications would vary, depending upon location and the availability of required materials. In South Carolina, where clay is plentiful and land is cheap, one small company has estimated it would cost approximately \$35,000 a year to capitalize and maintain a clay-lined pit for prospective use. In a more typical situation, an Illinois company estimated an annual capital cost of \$200,000 for an EPA-approved pit, without account for increased operating expenditures.

Construction of lined land fills, however, may not be feasible due to flood plain constraints, insufficient land capacity or the inability to meet all of the EPA specifications. In that case, recycling companies would have to find a chemically treated sanitary land fill which would accept their dross. Assuming that such land fills can be located (one plant in California identified such a facility, but was unable to transport the waste due to legal difficulties in crossing state lines), the costs here would be even more prohibitive. Transportation costs alone have been estimated at between \$3.50 and \$11.00 per ton, depending upon distance and mode of transfer; the cost of actually dumping the

dross would of course raise these figures substantially. In the Midwest, for example, plants have been advised of dumping costs at qualified facilities ranging between \$7 and \$30 per ton, exclusive of transportation. The only EPA-qualified land fill in South Carolina offered the recycler in that state a price of \$66 per ton, which calculated to a total disposal cost of \$900,000 a year; the quoted figure would have been more than enough to force the plant to close. Thus if the industry were forced to utilize chemically treated sanitary land fills exclusively, the disposal costs for 1970 alone, at South Carolina prices, could amount to as high as \$75 million.

Removal and disposal of presently stored wastes could add as much as an additional \$666 million, assuming that such land fill capacity could be located. Alternatively, the waste piles can be sealed, but the cost of such a disposal method would also be prohibitive. One plant in California considered having its 14 acre slag pile covered in concrete to avoid the annual maintenance cost. The lowest estimate was over \$1.25 million.

While the above costs by themselves are extraordinarily prohibitive for such large volumes of waste, cost data on complete compliance with all additional proposed hazardous waste regulations is not available at the present time. It is important to note, however, that these costs cannot be considered in a vacuum. secondary aluminum industry is composed of many small-sized companies dependent on low margin operations for their survival. Despite the recent emphasis on recycling and product conservation, few companies in the last five years have succeeded in making a profit, while most have not; some companies have failed to meet the margin and gone out of business altogether. The cost of compliance with EPA's proposed hazardous waste regulations and the introduction of expensive waste management systems would particularly burden the smaller-sized or marginal companies in the industry. Lacking the capitalization and financial stability of its sister companies in the primary aluminum industry, many ARA plants would simply be forced to stop operations. This conclusion is supported by EPA's own commissioned studies.

A 1973 study on the economic impact of similar proposed regulations, also prepared by the CALSPAN Corporation, concluded that "application of environmental regulations to the industry may increase capital requirements and minimum efficient size to such an extent that some smaller firms may exit from the industry, leading to increased concentration. The same regulations might also serve to raise the capital requirements barrier to entry into the secondary industry."

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Resource Conservation and Recovery Act, indicates that the presently proposed regulations would have a significantly adverse economic impact on the secondary aluminum industry. The study, prepared by Arthur D. Little, Inc., estimated that the annual cost of hazardous waste management could range from a low of over \$10 million to a high of approximately \$50 million. The ADL study concluded that industry compliance with the EPA hazardous waste program would result in either "probable" or "likely" plant closures, depending on the severity of the final regulations promulgated.

The evidence, as discussed in Section I, clearly does not justify the listing of the secondary aluminum industry as a generator of hazardous waste. The evidence becomes even more compelling when it is realized that compliance with the proposed regulations would force environmentally desirable companies out of business altogether. Thus the minimal benefits to be gained by subjecting an essentially clean industry to prohibitive regulation would be heavily outweighed by the substantial environmental benefits lost as a result of compliance.

The equities cry out for a more common sensible approach to the industry. Accordingly, the ARA respectfully requests that secondary aluminum dross smelting - high salt slag plant residue be taken off the list of hazardous wastes in any final rules promulgated.

APPENDIX



Summary of report of Herron Testing Laboratories, Inc. Cleveland, Ohio

Environmental Protection Agency (EPA) proposed regulations and guidelines are set forth in detail in the Federal Register of Monday, December 18, 1978 Part IV (Vol. 43, No. 243), pages 58946 thru 59028.

One of the wastes proposed to be hazardous is defined as "Secondary Aluminum Dross Smelting - High Salt Slag Plant Residue", SIC 3341. The basis for the proposed regulation to include this material depends on the amount of toxic elements that could be leached into the environment by natural processes. The proposed regulations list eight toxic metals. The toxic elements (Pg. 58956 in the cited Fed. Reg.) and their not to exceed concentrations are:

		1			Not	to e	xceed concen	tration
Element	Symbol			×	mg/1	l in	the extract	X 10
Arsenic	As		÷				0.50	
Barium	Ba				*		10.0	
Cadmium	Cd				8		0.10	•
Chromium	Cr	*		· · · · · · · · · · · · · · · · · · ·		90	0.50	
Lead	Pb	+					0.50	
Mercury	Hg				73		0.02	
Selemium	Se						0.10	
Silver	Ag						0.50	

Approved methodology as set forth in the Federal Register was followed.

Briefly, the extraction procedure consisted of treating 100 grams of slag with specific amounts of water and acctic acid for 24 hours within specific temperature limits (20° - 40° C). The resulting extract was then analyzed for the previously mentioned toxic elements by atomic absorption spectrophotometry (AA) and are expressed as milligrams per liter (mg/l) in the extract.

Five (5) member companies of the ARA shipped representative samples of their "High Salt Slag Residues" for evaluation.

The five (5) companies were designated by letters A thru E.

Each sample was analyzed at least twice, i.e., extracts were made on two (2) each 100 gram portions and are reported as run 1



and run 2. In some cases a third run was made and is reported as run 3.

One company (E) when apprised of their slag analysis submitted another sample (E-2) which is a slag from the same process but deleting non-magnetic fines of auto-shredder scrap. Further the dross sample submitted was crushed to -20 mesh, approximately the same particle size as the samples submitted by Plants A thru D.

The analytical data is attached.

Company		Α		В		. (<u>(1)</u>	«.	p`	
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	7.		30						38 12	
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					•	<u>L 7</u>			L 7194	
Run No.	<u>#1</u>		<u>#1</u>	#2	<u>∜3</u>	. 71	72	<u>#1</u>	<u>#2</u>	<u>#3</u>
Element Max								w.		
parentheses								•		7
<u> 55/1 X 10</u>	•	7.		592						
		20					ž	¥	E " 8	
Ag (.50)	0.02	0.04	0.08	0.01		0.05	0.08	<0.01	<0.01	
Ba (10.0)	5.5	5.1	1.2	1.7	9 . %	4.7	3.0	1.0	2.3	
Cd (0.10)	<0.05	<0.05	<0.05	<0.05		<0.05	<0.05	0.03	0.07	
As (0.50)	<0.001	0.002	. 0.001	0.001		0.002	0.001	<0.001	<0.001	
Hg (0.02)	<0.0002	0.0002	0.000	4 0.0003		0.0004	0.0003	0.0004	0.0007	
Se (0.10)	<0.01 <	0.01	<0.01	<0.01		<0.01	<0.01.	<0.01	<0.01	
Cr (0.50)	0.07	0.07	* 0.04	0.08		0.02	0.07	<0.02	0.08	φ ¹ α
Pb (0.50)	0.05	0.06	0.06	0.45	0.28	<0.05	0.10	<0.05	0.60	0.09
							e:		23	
< - less than			3				E	~ ×	1	1
•				9				Dn.n.l.n 3/1./7		e.y

E-2

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	Pò		2.8	4.8	w ⁼ v	0.11	0.08

< - less than '

* - not complete at reporting time. To be furnished at later date.

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Section 13

(Red) November 18, 1975

Chief

SUBJECT

Request for Policy Decision Direct Discharge to Ground Water U. S. Aluminum Corporation Marietta Borough, Lancaster County

TO

Gary L. Merritt, Chief Technical Services Unit Ground Water Section

FROM

The state of the s

Jeff Peffer Regional Geologist Harrisburg Region Thru: James T. Flesher, Operations Section

The above industry generates 20,000 GPD of waste water which has high concentrations of dissolved solids, most of which is potassium chloride. This waste water is currently discharged to an infiltration basin at the east end of the plant complex, on the flood plain of the Susquehanns River. The writer made a field inspection of the area on 11/14/75.

The basin lies approximately 350'north of the river, and is "pinpointed" on the attached copy of a portion of the Columbia West 7½ minute quadrangle. The area is immediately underlain by an undetermined thickness of sandy alluvium. In a gully along a road to the east of the lagoon, orangish-brown sandy silt is exposed. Bedrock mapped as underlying the area is argillaceous dolomite of the Vintage formation. No bedrock outcrops were observed on the date of the writer's field

Ground water in the site area is expected to be flowing to the south, towards and discharging to the river. Depending on the nature of the underlying flow system(s) and the depth of alluvium, ground water may have components of flow more or less parallel to the river, along bedrock structure.

At this point, in order for U. S. Aluminum Corporation to receive a permit to discharge this waste to the river, the basin would have to be lined (to meet the impermeability requirement of Chapter 101 of the Rules and Regulations) and a dispersion device (required by Facilities Section) would have to be provided at the discharge point(s).

The applicant's engineer has rightfully questioned whether the existing infiltration basin - ground water mixing zone scheme would not be less detrimental to the environment than a concentrated surface discharge.

In a meeting with the engineer I indicated that concentrated direct discharges to ground water are not in line with current Departmental policy. However, I intimated that, with the concurrence of Central Office Ground Water Section, such a discharge might be permitted (as a direct discharge and not a lagoon) if:

